Refine Search

Search Results -

Terms	Documents	
L11 same L3	32	

US Pre-Grant Publication Full-Text Database
US Patents Full-Text Database
US OCR Full-Text Database
EPO Abstracts Database

Database:

EPO Abstracts Database
JPO Abstracts Database
Derwent World Patents Index
IBM Technical Disclosure Bulletins

Search:

12	[4]
	4 10000F
	# P0000000 900000000 900000000
	1824

Refine Search





Interrupt

Search History

DATE: Monday, March 01, 2004 Printable Copy Create Case

Set Name Query side by side		Hit Count Set Name result set		
DB=USPT; PLUR=YES; OP=ADJ				
	<u>L12</u>	L11 same 13	32	<u>L12</u>
	<u>L11</u>	il-6 same (11 or 15) same 17	95	<u>L11</u>
	<u>L10</u>	L9 and 18	48	<u>L10</u>
	<u>L9</u>	il-6	4419	<u>L9</u>
	<u>L8</u>	L7 and 16	69	<u>L8</u>
	<u>L7</u>	bone	66866	<u>L7</u>
	<u>L6</u>	L5 same 13	102	<u>L6</u>
	<u>L5</u>	anti-cytokine	207	<u>L5</u>
	<u>L4</u>	L3 same 12 same 11	2074	<u>L4</u>
	<u>L3</u>	in vivo or therapy or inhibit\$	422134	<u>L3</u>
	<u>L2</u>	cytokine or cytokines	17324	<u>L2</u>
	<u>L1</u>	antibody or antibodies	67564	<u>L1</u>

END OF SEARCH HISTORY

```
4994 S IL-11 OR INTERLEUKIN-11
            673 S L1 (P) ANTIBOD?
1.2
            404 S L2 (P) (ADMINIST? OR THERAP? OR TREAT? OR METHOD?)
T.3
=> s l1 (5a) antibod?
            89 L1 (5A) ANTIBOD?
L4
=> s 14 (5a) (administ? or therap? or treat? or method?)
             9 L4 (5A) (ADMINIST? OR THERAP? OR TREAT? OR METHOD?)
L_5
=> d 15 1-9 bib kwic
     ANSWER 1 OF 9
                       MEDLINE on STN
1.5
     2002322722
                    MEDLINE
AN
     PubMed ID: 12065535
DN
TI
     Contribution of interleukin-11 and prostaglandin(s) in
     lipopolysaccharide-induced bone resorption in vivo.
     Li Li; Khansari Alireza; Shapira Lior; Graves Dana T; Amar Salomon
ΑU
     Department of Periodontology and Oral Biology, School of Dental Medicine,
CS
     Boston University, Boston, Massachusetts 02118, USA.
     12482 (NIDCR)
NC
     DE07559
     Infection and immunity, (2002 Jul) 70 (7) 3915-22.
SO
     Journal code: 0246127. ISSN: 0019-9567.
CY
     United States
     Journal; Article; (JOURNAL ARTICLE)
DT
LΑ
     English
FS
     Priority Journals
EM
     200207
ED
     Entered STN: 20020615
     Last Updated on STN: 20020731
     Entered Medline: 20020730
AB
     . . . lacking IL-1 receptor type I (IL-1RI(-/-)), mice lacking TNF
     receptor p55 and IL-1RI (TNFRp55(-/-)-IL-1RI(-/-)), and wild-type mice.
     Mice were then treated with injections of anti-IL-
     11 monoclonal antibody (MAb), indomethacin, or
     phosphate-buffered saline (PBS) and sacrificed 5 days later. Histological
     sections stained for tartrate-resistant acid phosphatase (TRAP) were.
     ANSWER 2 OF 9
                       MEDLINE on STN
L5
ΑN
     1999069250
                    MEDLINE
DN
                PubMed ID: 9767454
     An anti-inflammatory role for interleukin-11 in established murine
     collagen-induced arthritis.
     Walmsley M; Butler D M; Marinova-Mutafchieva L; Feldmann M
ΑU
     Kennedy Institute of Rheumatology, London, UK.
CS
     IMMUNOLOGY, (1998 Sep) 95 (1) 31-7.
SO
     Journal code: 0374672. ISSN: 0019-2805.
     ENGLAND: United Kingdom
CY
DT
     Journal; Article; (JOURNAL ARTICLE)
     English
LΑ
FS
     Priority Journals
EM
     199901
ED
     Entered STN: 19990209
     Last Updated on STN: 19990209
     Entered Medline: 19990128
     . . of IL-11 that the anticollagen type II (CII) response may have
AB
     been augmented, there was no statistically significant effect of
     IL-11 treatment on anti-CII antibody
     levels. Similarly, the acute-phase reactant serum amyloid P was only
     elevated in mice receiving very high doses (50-100 microgram/day) of.
```

```
ANSWER 3 OF 9 CAPLUS COPYRIGHT 2004 ACS on STN
L5
AN
     2002:965157 CAPLUS
DN
     138:38076
     WSX receptor agonist antibodies for treating diseases associated with
TI
     lymphopoiesis, erythropoiesis, myelopoiesis, and obesity
     Carter, Paul J.; Chiang, Nancy Y.; Kim, Kyung Jin; Matthews, William;
IN
     Rodrigues, Maria L.
DΔ
     U.S. Pat. Appl. Publ., 140 pp., Cont.-in-part of U.S. Ser. No. 667,197.
SO
     CODEN: USXXCO
DT
     Patent
LA
     English
FAN.CNT 2
     PATENT NO.
                                           APPLICATION NO.
                  KIND
                            DATE
                                           US 1997-779457
     US 2002193571
                            20021219
                                                             19970107
PI
                      A1
     CA 2241564
                                           CA 1997-2241564
                      AA
                            19970717
                                                            19970107
     ZA 9700148
                      Α
                            19980708
                                           ZA 1997-148
                                                            19970108
     US 6541604
                                           US 1997-780562
                       B1
                            20030401
                                                            19970108
                                           US 2002-214802
     US 2003004109
                      A1
                            20030102
                                                            20020806
PRAI US 1996-585005
                       B2
                            19960108
     US 1996-667197
                       Α2
                            19960620
     US 1996-64855P
                       Ρ
                            19960108
     US 1997-780562
                       A3
                            19970108
IT
     Cytokines
     Interleukin 1
     Interleukin 10
       Interleukin 11
     Interleukin 2
     Interleukin 3
     Interleukin 4
     Interleukin 5
     Interleukin 6
     Interleukin 7
     Interleukin 8
     Interleukin 9
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (WSX receptor antibodies for treating diseases
        associated with lymphopoiesis, erythropoiesis, myelopoiesis, and obesity)
L_5
     ANSWER 4 OF 9 CAPLUS COPYRIGHT 2004 ACS on STN
AN
     2002:489736 CAPLUS
DN
     137:183745
TI
     Contribution of interleukin-11 and prostaglandin(s) in
     lipopolysaccharide-induced bone resorption in vivo
ΑU
     Li, Li; Khansari, Alireza; Shapira, Lior; Graves, Dana T.; Amar, Salomon
CS
     Department of Periodontology and Oral Biology, School of Dental Medicine,
     Boston University, Boston, MA, 02118, USA
SO
     Infection and Immunity (2002), 70(7), 3915-3922
     CODEN: INFIBR; ISSN: 0019-9567
PB
     American Society for Microbiology
DT
     Journal
     English
Τλ
RE.CNT 41
              THERE ARE 41 CITED REFERENCES AVAILABLE FOR THIS RECORD
              ALL CITATIONS AVAILABLE IN THE RE FORMAT
     We previously demonstrated that interleukin-1 (IL-1) and tumor necrosis
     factor (TNF) activities only partially account for calvarial bone
     resorption induced by local application of lipopolysaccharide (LPS) in
            The present study was undertaken to determine the role and relative
     contribution of IL-11 and prostaglandin(s) (PG[s]) in LPS-induced bone
     resorption in vivo. A one-time dose of LPS was injected into the s.c.
     tissue overlying calvaria of mice lacking IL-1 receptor type I
     (IL-1RI-/-), mice lacking TNF receptor p55 and IL-1RI (TNFRp55-/--IL-1RI-/-
     ), and wild-type mice. Mice were then treated with injections
```

of anti-IL-11 monoclonal antibody (MAb), indomethacin, or phosphate-buffered saline (PBS) and sacrificed 5 days later. Histol. sections stained for tartrate-resistant acid phosphatase (TRAP) were quantified by histomorphometric anal. At low doses of LPS (100 µg/mouse), the percentages of bone surface covered by osteoclasts were found to be similar in three strains of mice. The increase was reduced by 37% with anti-IL-11 MAb and by 46% with indomethacin. At higher doses of LPS (500 $\mu g/mouse$), we found an eightfold increase in these percentages in wild-type mice and a fivefold increase in these percentages in IL-1RI-/- and TNFRp55-/--IL-1RI-/- mice after normalizing with the value from the saline-PBS control group in the same strain of mice. The increase was reduced by 55 and 69% in wild-type mice and by 50 and 57% in IL-1RI-/- and TNFRp55-/--IL-1RI-/- mice treated with anti-IL-11 MAb or indomethacin, resp. Our findings suggest that in vivo, at low doses of LPS (100 µg/mouse), LPS-induced bone resorption is mediated by IL-11 and PGs, while at high doses of LPS (500 $\mu g/mouse$), it is mediated by IL-11, PGs, IL-1, and TNF signaling. IL-11 and PGs mediate LPS-induced bone resorption by enhancing osteoclastogenesis independently of the IL-1 or TNF signaling. ANSWER 5 OF 9 CAPLUS COPYRIGHT 2004 ACS on STN 2001:101180 CAPLUS 134:161896 Mammalian cytokines; related reagents Timans, Jacqueline C.; Kastelein, Robert A.; Bazan, J. Fernando Schering Corporation, USA PCT Int. Appl., 73 pp. CODEN: PIXXD2 Patent English FAN.CNT 1 PATENT NO. KIND DATE APPLICATION NO. DATE ____ _____ ______ WO 2001009176 A2 20010208 WO 2000-US20475 20000727 A3 WO 2001009176 20011101 W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, HR, HU, ID, IL, IN, IS, JP, KG, KR, KZ, LC, LK, LR, LT, LU, LV, MA, MD, MG, MK, MN, MX, MZ, NO, NZ, PL, PT, RO, RU, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UZ, VN, YU, ZA, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG EP 1200592 A2 20020502 EP 2000-950787 20000727 AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL T2JP 2001-513982 JP 2003506026 20030218 20000727 PRAI US 1999-364674 A 19990730 US 1999-369643 Α 19990806 WO 2000-US20475 W 20000727 Antibodies RL: BSU (Biological study, unclassified); BUU (Biological use, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (mammalian interleukin 80D and interleukin 11 for diagnosis and therapy of degenerative or abnormal condition of immune system and/or hematopoietic cells) Antibodies RL: BSU (Biological study, unclassified); BUU (Biological use, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (monoclonal; mammalian interleukin 80D and interleukin

11 for diagnosis and therapy of degenerative or

abnormal condition of immune system and/or hematopoietic cells)

L5

AN

DN

TI

IN

PA

SO

DT

LA

РΤ

TT

IT

```
AN
     1999:753092 CAPLUS
DN
     132:2795
TI
     Antagonists of interleukin 11-mediated osteoporotic bone loss
IN
     Shaughnessy, Stephen; Austin, Richard Carl
PΑ
     Hamilton Civic Hospital Research Development Corporation, Can.
     PCT Int. Appl., 61 pp.
SO
     CODEN: PIXXD2
DT
     Patent
LΑ
     English
FAN.CNT 1
     PATENT NO.
                      KIND
                            DATE
                                           APPLICATION NO.
                                                             DATE
PΙ
     WO 9959608
                       A2
                            19991125
                                           WO 1999-CA516
                                                             19990519
     WO 9959608
                       A3
                            20000406
            AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ,
             DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS,
             JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK,
             MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ,
             TM, TR, TT, UA, UG, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD,
             RU, TJ, TM
         RW: GH, GM, KE, LS, MW, SD, SL, SZ, UG, ZW, AT, BE, CH, CY, DE, DK,
             ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG,
             CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
     CA 2328486
                       AA
                            19991125
                                           CA 1999-2328486
                                                             19990519
     AU 9940277
                       A1
                            19991206
                                           AU 1999-40277
                                                             19990519
     AU 767749
                            20031120
     EP 1079847
                       A2
                            20010307
                                           EP 1999-923352
                                                             19990519
            AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
             IE, FI
     JP 2002515444
                       T2
                            20020528
                                           JP 2000-549272
                                                             19990519
PRAI CA 1998-2237915
                            19980519
                       Α
     WO 1999-CA516
                            19990519
IT
     Antibodies
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (neutralizing; to interleukin-11 or IL-11 receptor
        for osteoporosis therapy)
L5
     ANSWER 7 OF 9 CAPLUS COPYRIGHT 2004 ACS on STN
AN
     1998:586896
                 CAPLUS
DN
     129:301592
TΤ
     An anti-inflammatory role for interleukin-11 in established murine
     collagen-induced arthritis
ΑU
     Walmsley, M.; Butler, D. M.; Marinova-Mutafchieva, L.; Feldmann, M.
CS
     Kennedy Inst. of Rheumatology, London, UK
SO
     Immunology (1998), 95(1), 31-37
     CODEN: IMMUAM; ISSN: 0019-2805
PΒ
     Blackwell Science Ltd.
     Journal
DT
LA
     English
RE.CNT
       31
              THERE ARE 31 CITED REFERENCES AVAILABLE FOR THIS RECORD
              ALL CITATIONS AVAILABLE IN THE RE FORMAT
AB
     Interleukin-11 (IL-11) is a cytokine belonging to the IL-6 family which
    has both pro- and anti-inflammatory potential. Like IL-6 it can diminish
     tumor necrosis factor-\alpha and IL-1 production, and augment Ig synthesis.
    We have explored the immunomodulatory effects of IL-11 treatment in mice
     in a model of inflammatory autoimmune joint disease, collagen-induced
     arthritis (CIA). Recombinant human IL-11 was administered at various
    doses to DBA/1 mice after the onset of CIA. IL-11 treatment caused a
    significant reduction in the clin. severity established CIA, which was
associated
    with protection from joint damage, as assessed by histol. Although there
    was a suggestion at high doses of IL-11 that the anticollagen type II
```

(CII) response may have been augmented, there was no statistically

significant effect of IL-11 treatment on

anti-CII antibody levels. Similarly, the acute-phase reactant serum amyloid P was only elevated in mice receiving very high doses (50-100 $\mu g/day$) of IL-11. Endogenous IL-11 was abundantly produced in synovial membrane cultures derived from CII-immunized mice with active disease, suggesting that, as in rheumatoid arthritis, this cytokine is spontaneously produced in the inflammatory response in CIA. The results presented here demonstrate an anti-arthritic immunoregulatory role for IL-11 in murine CIA, and suggest that IL-11 is a candidate therapeutic

L5

ΑN

DN

TI

IN

PA

SO

DT

LA

PΙ

IT

1.5

ΑN

TΤ

TN

PΑ

PΤ

ΑI

DТ

FS

ECL

DETD

```
mol. for human inflammatory arthritic diseases.
     ANSWER 8 OF 9 CAPLUS COPYRIGHT 2004 ACS on STN
     1996:681971 CAPLUS
     126:6450
     Anti-gp130 monoclonal antibodies for inflammatory disease treatment
     Burstein, Samuel A.
     The Board of Regents of the University of Oklahoma, USA
     U.S., 14 pp.
     CODEN: USXXAM
     Patent
     English
FAN.CNT 1
     PATENT NO.
                      KIND DATE
                                           APPLICATION NO.
                                                            DATE
                                           -----
     US 5571513
                      Α
                            19961105
                                           US 1995-455799
                                                            19950531
     WO 9638481
                      A1
                            19961205
                                           WO 1996-US7385
                                                            19960522
         W: AU, CA, JP
         RW: AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE
     AU 9658695
                       A1
                            19961218
                                           AU 1996-58695
                                                            19960522
PRAI US 1995-455799
                            19950531
     WO 1996-US7385
                            19960522
     Interleukin 11
     Interleukin 6
     RL: BAC (Biological activity or effector, except adverse); BPR (Biological
     process); BSU (Biological study, unclassified); BIOL (Biological study);
     PROC (Process)
        (anti-gp130 monoclonal antibodies for inflammatory disease
        treatment)
     ANSWER 9 OF 9 USPATFULL on STN
       2002:19059 USPATFULL
       Agonist antibodies
       Adams, Camellia W., Mountain View, CA, United States
       Carter, Paul J., San Francisco, CA, United States
       Fendly, Brian M., Half Moon Bay, CA, United States
       Gurney, Austin L., Belmont, CA, United States
       Genentech, Inc., South San Francisco, CA, United States (U.S.
       corporation)
       US 6342220
                          В1
                               20020129
       US 1997-918148
                               19970825 (8)
       Utility
       GRANTED
EXNAM
      Primary Examiner: Nolan, Patrick J.
LREP
       Piper Marbury Rudnick & Wolfe LLP, Kelber, Steven B.
CLMN
      Number of Claims: 14
       Exemplary Claim: 1
       11 Drawing Figure(s); 11 Drawing Page(s)
DRWN
LN.CNT 4209
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       . . . stimulating factors or interleukins include; kit-ligand, LIF,
      G-CSF, GM-CSF, M-CSF, EPO, IL-1, IL-2, IL-3, IL-5, IL-6, IL-7, IL-8,
       IL-9 or IL-11. Alternatively, the antibody
       is administered in combination with an Insulin-like growth
      factor (e.g., IGF-1) or a tumor necrosis factor (e.g., lymphotoxin
       (LT)).
```